attach paper no. 7

PATENT Our Docket: P31 8756

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application f: TULLIS

Serial No.: 07/633,453

Filed: December 20, 1990

For: OLIGONUCLEOTIDE THERAPEUTIC AGENT AND <u>NETHODS OF MAKING SAME</u> Group Art Unit: 185

Examiner: J. Martinell

BEST AVAILABLE COPY

Hon. Commissioner of Patents and Trademarks Washington, D.C. 20231

COMMUNICATION

sir:

The Examiner requested a proposed amendment be forwarded for his review prior to the interview scheduled with Applicant's attorney, Cathryn Campbell, during the week of October 14, 1991. Therefore, transmitted herewith is a proposed amendment to the Office Action mailed on August 6, 1991, for the above-identified application.

Respectfully submitted,

Theresa A. Brown

Registration No. 32,547 Telephone: (619) 535-9001 Pacsimile: (619) 535-8949

PRETTY, SCHROEDER, BRUEGGEMANN & CLARK 444 South Plower Street Suite 2000 Los Angeles, California CERTIFICATE OF FACSIMILE TRANSMISSION

I hereby curtify that these 4 papers are being facsimile transmitted to the Patent and Trademark Office on date shown below:

STEPHEN R. REITER

19/10/91

OCT 10 1991

DIRECTUR GRUUr

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DRAFT

PATENT Our Docket: P31 8756

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application f: TULLIS

4524

Serial No.: 07/633,453

Filed: December 20, 1990

For: OLIGONUCLEOTIDE

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Examiner: J. Martinell

Hon. Commissioner of Patents and Trademarks Washington, D.C. 20231

PROPOSED RESPONSE TO OFFICE ACTION

Sir:

In response to the Office Action mailed on August 6, 1991, Applicant requests entry of the amendments and consideration of the remarks set forth below.

PROPOSED AMENDMENTS

In the Claims:

53. (amended) A method of developing oligodeoxyribonucleotide therapeutic agents for use in in vivo inhibition of the synthesis of one or more targeted proteins in a cell without substantially inhibiting the synthesis of non-targeted proteins, comprising the steps of:

determining the base sequence of an organism's messenger ribonucleic acid, said base sequence coding for at least a portion of said protein targeted for inhibition;

synthesizing an oligodeoxyribonucleotide, the nucleotide sequence of which is substantially complementary to at least a portion of said base sequence and capable of hybridization with said messenger ribonucleic acid base sequence